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WHAT IS CLAIMED IS:

1. A molecule comprising a polypeptide having substantial homology with a CTL epitope selected from the 5 group consisting of ADLMGYIPLV (Core₁₃₁₋₁₄₀; SEQ ID NO:1), LLALLSCLTV (Core₁₇₈₋₁₈₇; SEQ ID NO:2), QLRRHIDLLV (SEQ ID NO:55), LLCPAGHAV (NS3₁₁₆₉₋₁₁₇₇; SEQ ID NO:26), KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅; SEQ ID NO:28), SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV (NS4₁₈₀₇₋₁₈₁₆; SEQ ID NO:35), and ILDSFDPLV (NS5₂₂₅₂₋₂₂₆₀; SEQ ID NO:42).

- 2. The molecule of claim 1, wherein said molecule comprises at least about eight amino acids and less than about 50 amino acids.
- 3. The molecule of claim/2, wherein said molecule comprises at least about nine amino acids and less than about thirteen amino acids.
- 20 4. The molecule of claim 1, wherein said polypeptide is KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅; SEQ ID NO:28) or substantially homologous thereto.
- 5. The molecule of claim 1, wherein said polypeptide is conjugated to a substance, wherein said substance is selected from the group consisting of a radiolabel, an enzyme, a fluorescent label, a solid matrix, a carrier, and a second CTL epitope.
- 30 6. The molecule of claim 5, wherein said substance is a second CTL epitope.
 - 7. The molecule of claim 5, wherein said second epitope is a T helper epitope.

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- The molecule of claim 5, wherein said carrier comprises an immunogenic lipid or protein.
- The molecule of claim 5, wherein said 9. polypeptide is conjugated to said substance indirectly by 5 a linker.
 - (10). A polypeptide having substantial homology with a CTE epitope selected from the group consisting of ADLMGYIPLV (Core₁₃₁₋₁₄₀; SEQ ID NO:1), LLALLS¢LTV (Core₁₇₈₋₁₈₇; SEQ ID NO:2), QLRRHIDLLV (SEQ ID NO:55), /LLCPAGHAV (NS31169- $_{1177}$; SEQ \(\text{TD NO:26} \) , KLVALGINAV (NS3 $_{1406-1415}$; /SEQ ID NO:28) , SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV (NS4₁₈₀₇₋₁₈₁₆; SEQ ID NO $\{35\}$, and ILDSFDPLV (NS5₂₂₅₂₋₂₂₆₀/; SEQ ID NO:42).
 - The polypeptide of claim 10, wherein said polypeptide is KLVALGINAV (NS3_{1406-14/5}; SEQ ID NO:28) or substantially homologous thereto,
- A method of provoking an immune response to a hepatitis C viral antigen, comprising contacting a cytotoxic T (ymphocyte/with/an immune response provoking amount of a molecule domprising a peptide selected from the group consisting of ADLMGYIPLV (Core,31-140; SEQ ID NO:1), LLALLSCLTV (Core,78-187; SEQ ID NO:2), QLRRHIDLLV (SEQ 25 ID NO:55), LLCPACHAV MS31169-1177; SEQ ID NO:26), KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅; SEQ ID NO: 28), SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV $\sqrt{(NS4_{1807-1816}; SEQ ID NO:35)}$, and ILDSFDPLV $(NS5_{2252-2260}; SEQ ID /NQ:42).$
 - The method of claim 12, wherein said contacting occurs in a mammal.
- The method of claim 13, wherein said mammal is 35 free of ACV disease, is a carrier of HCV, or is afflicted with HQV disease.

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The method of claim 12, wherein said contacting occurs in vitro.

- The method of claim 15, wherein said method further comprises returning said contacted cytotoxic T cells to the host.
- The method of claim 12, wherein said polypeptide is co-administered with a second polypeptide that induces a T helper response to HCV.
 - 18) The method of claim 17, wherein said polypeptide and said T helper inducing polypeptide are conjugated to one another.

Amethod of detecting in lymphocytes of a mammal cytotoxic T cells that respond to a T cell epitope of hepatitist virus, comprising the steps of: (a) contacting target cells with a molecule comprising at least one of the peptides selected from the group 20 consisting of ADLMGYIPLV (Core 131-140; SEQ ID NO:1), LLALLSCLTV (dore 178-187; SEQ /D NO:2), QLRRHIDLLV (SEQ ID NO:55), LLCPAGHAV (NS311/09-1/17; SEQ ID NO:26), KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅; SEQ ID NO: 28) / SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV $\sqrt{N_{5}^{4}4_{1807-1816}}$; SEQ ID NO:35), and ILDSFDPLV 25 (NS52252-2260; SEQ ID MO:42), wherein said target cells are of the same HLA class as the lymphocytes to be tested for said cytotoxic T cells; (b) contacting said lymphocytes to be tested for said cytotoxic T cells with a molecule comprising at least one of the peptides selected from the 30 peptides selected from the group consisting of ADLMGYIPLV (Core₁₃₁₋₁₄₀; SEØ ID NO:1), LLALLSCLTV (Core₁₇₈₋₁₈₇; SEQ ID NO:2), LLCPÁGHA∜ (NS3₁₁₆₉₋₁₁₇₇; SEQ ID NO:26), KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅;/SEQ ID NO:28), SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV (NS4₁₈₀₇₋₁₈₁₆; SEQ ID NO:35), and ILDSFDPLV 35 (NS5₂₂₅₂/₂₂₆₀; SEQ ID NO:42); and (c) determining whether said

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lymphocytes exert a cytotoxic effect on said target

20. A pharmaceutical composition comprising a molecule that includes a polypeptide having substantial homology with a CTL epitope selected from the group consisting of ADLMGYIPLV (Core₁₃₁₋₁₄₀; SEQ ID/NO:1), LLALLSCLTV (Core₁₇₈₋₁₈₇; SEQ ID NO:2), QLRRHIDLLV (SEQ ID NO:55), LLCPAGHAV (NS3₁₁₆₉₋₁₁₇₇; SEQ ID NO:26), KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅; SEQ ID NO:28), SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV (NS4₁₈₀₇₋₁₈₁₆; SEQ ID NO:35), and ILDSFDPLV (NS5₂₂₅₂₋₂₂₆₀; SEQ ID NO:42), and a pharmaceutically acceptable carrier.

A pharmaceutical composition comprising a polypeptide having substantial homology with a CTL epitope selected from the group consisting of ADLMGYIPLV (Core₁₃₁₋₁₄₀; SEQ ID NO:1), LLALLSCLTV (Core₁₇₈₋₁₈₇; SEQ ID NO:2), OLRRHIDLLV (SEQ ID NO:55), LLCPAGHAV (NS3₁₁₆₉₋₁₁₇₇; SEQ ID NO:26, KLVALGINAV (NS3₁₄₀₆₋₁₄₁₅; SEQ ID NO:28), SLMAFTAAV (NS4₁₇₈₉₋₁₇₉₇; SEQ ID NO:34), LLFNILGGWV (NS4₁₈₀₇₋₁₈₁₆; SEQ ID NO:35), and ILDSFDPLV (NS5₂₂₅₂₋₂₂₆₀; SEQ ID NO:42), and a pharmaceutically acceptable carrier.

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